## In the specification.

The specification is amended to correctly recite the priority of this application. No new matter is added by this Amendment.

If a telephone conference would expedite prosecution of this application, the Examiner is invited to telephone the undersigned at (415) 217-6022.

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Respectfully submitted,

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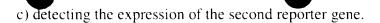
## **CLAIMS PENDING AS OF DECEMBER 4, 2000**

- A method for screening a test compound for the ability to activate transcription through an indirect estrogen response, the method comprising:
- a) providing a cell comprising an estrogen receptor and a promoter comprising an AP1 site which regulates expression of a reporter gene;
  - b) contacting the cell with the test compound; and
  - c) detecting the expression of the reporter gene.
  - 2. (Amended) A method of claim 1, wherein the cell is an Ishikawa cell.
- 3. (Amended) A method of claim 1, wherein the cell over-expresses the estrogen receptor.
- 4. (Amended) The method of claim 1, wherein the promoter is genetically engineered to comprise an AP1 site.
- 5. (Amended) The method of claim 1, wherein the test compound is known to have antiestrogenic activity.
- 6. (Amended) The method of claim 1, wherein the cell is derived from uterine tissue.
- 7. (Amended) The method of claim 6, wherein the cell is a HeLa cell or an Ishikawa cell.
  - 8. (Amended) A method of claim 1, further comprising the steps of:

comprising a standard estrogen response element which regulates expression of a second reporter gene;

b) contacting the second cell with the test compound, and

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- 9. (Amended) A method of claim 8, wherein the response element is from the *Xenopus* vitellogenin A2 gene.
- 10. (Amended) A method of claim 1, wherein the cell further comprises a promoter comprising a standard estrogen response element which regulates expression of second reporter gene.
- 11. (Amended) A method of claim 10, wherein the response element is from the *Xenopus* vitellogenin A2 gene.
  - 12. (Amended) An estrogen agonist identified by the method of claim 1.
- A method for screening a test compound for the ability to inhibit transcription through an indirect estrogen response, the method comprising:
- a) providing a cell comprising an estrogen receptor and a promoter comprising an AP1 site which regulates expression of a reporter gene;
- b) contacting the cell with the test compound an a compound known to mediate an indirect estrogen response;
  - c) detecting the expression of the reporter gene.
- 14. (Amended) The method of claim 13, wherein the compound is known to mediate an indirect estrogen response is tamoxifen.
- 15. (Amended) A method of claim 13, wherein the cell over-expresses the estrogen receptor.
- 16. (Amended) The method of claim 13, wherein the promoter is genetically engineered to comprise an AP1 site.

Amended - a compound identified by the method of claim 13.

A method for screening a test environmental compound for estrogenic activity, the method comprising:

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- a) providing a cell comprising an estrogen receptor and a promoter comprising an estrogen response element which regulates the expression of a reporter gene;
  - b) contacting the cell with the test compound; and
  - c) detecting the expression of the reporter gene.
- 19. (Amended) The method of claim 18, wherein the cell further comprises a promoter comprising an AP1 site which regulates expression of a second reporter gene.
  - 20. (Amended) The method of claim 18, wherein the reporter gene is CAT.
- 21. (Amended) The method of claim 18, wherein the cell over-expresses the estrogen receptor.
  - 22. (Amended) The method of claim 18, wherein the cell is an ERC1 cell.
- A method of inhibiting agonistic activity of an antiestrogen compound, said method comprising administering with said antiestrogen compound an inhibitor selected from the group consisting of genistein, staurosporine, 6-thioguanine, and 2 aminopurine.
- 24. (Amended) The method of claim 23, wherein said inhibiting agonistic activity comprises inhibiting an indirect estrogen response.
- 25. (Amended) The method of claim 23, wherein said antiestrogen compound is tamoxifen.
  - 26. (Amended) The method of claim 23, wherein said inhibition is *in vivo*.

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